



The
Patent
Office

09/83

PCT/GB 99 / 04099

24 DECEMBER 1999

INVESTOR IN PEOPLE

GB99/4099

PRIORITY DOCUMENT

SUBMITTED OR TRANSMITTED IN
COMPLIANCE WITH RULE 17.1(a) OR (b)

The Patent Office
Concept House
Cardiff Road
Newport
South Wales

NP10 8QQ	
REC'D 13 JAN 2000	
WIPO	PCT

I, the undersigned, being an officer duly authorised in accordance with Section 74(1) and (4) of the Deregulation & Contracting Out Act 1994, to sign and issue certificates on behalf of the Comptroller-General, hereby certify that annexed hereto is a true copy of the documents as originally filed in connection with the patent application identified therein.

In accordance with the Patents (Companies Re-registration) Rules 1982, if a company named in this certificate and any accompanying documents has re-registered under the Companies Act 1980 with the same name as that with which it was registered immediately before re-registration save for the substitution as, or inclusion as, the last part of the name of the words "public limited company" or their equivalents in Welsh, references to the name of the company in this certificate and any accompanying documents shall be treated as references to the name with which it is so re-registered.

In accordance with the rules, the words "public limited company" may be replaced by p.l.c., plc, P.L.C. or PLC.

Re-registration under the Companies Act does not constitute a new legal entity but merely subjects the company to certain additional company law rules.

Signed

Dated

16 DEC 1999

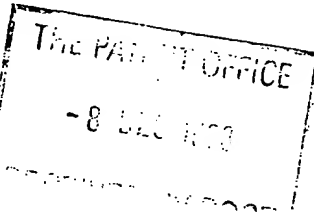
926845.1

1 ne.

Ac. 77

DEC 1998

98DEC98 E410771-1 002776
P01/7700 0.00 - 9826845.1



Request for grant of a patent

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)

The Patent Office

Cardiff Road
Newport
Gwent NP9 1RH

1. Your reference

DERA/IPD01/P2746

2. Patent application number

(The Patent Office will fill in this part)

- 8 DEC 1998

3. Full name, address and postcode of the or of each applicant (underline all surnames)

THE SECRETARY OF STATE FOR DEFENCE
DEFENCE AND EVALUATION RESEARCH AGENCY
IVELY ROAD
FARNBOROUGH GU14 OLX

Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

UNITED KINGDOM

07349496001

4. Title of the invention

ORGANIC LIGHT EMITTING MATERIAL AND DEVICE

5. Name of your agent (if you have one)

BOWDERY ANTHONY OLIVER

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

BOWDERY ANTHONY OLIVER
D/IPR FORMALITIES SECTION (DERA)
POPLAR 2
MOD(PE) ABBEY WOOD #19
BRISTOL
BS34 8JH

Patents ADP number (if you know it)

06935910003

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country

Priority application number
(if you know it)

Date of filing
(day / month / year)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number or earlier application

Date of filing
(day / month / year)

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:

YES

- a) any applicant named in part 3 is not an inventor, or
- b) there is an inventor who is not named as an applicant, or
- c) any named applicant is a corporate body.

See note (d))

9. Enter the number of sheets for any of the following items you are filing with this form. Do not count copies of the same document

Continuation sheets of this form

Description 13 /
 Claim(s) 2 /
 Abstract 1 /
 Drawing(s) 4 *4-10*

10. If you are also filing any of the following, state how many against each item.

Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (*Patents Form 7/77*)

1 + 2

Request for preliminary examination and search (*Patents Form 9/77*)

1

Request for substantive examination (*Patents Form 10/77*)

Any other documents
(*please specify*)

11. I / We request the grant of a patent on the basis of this application.

Signature

J.B. EDWARDS

Date 3/12/98

12. Name and daytime telephone number of person to contact in the United Kingdom MR J.B.EDWARDS 01684 894002

Warning

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent of the same invention and either no direction prohibiting publication or communication has been given, or any such direction has been revoked.

Notes

- If you need help to fill in this form or have any questions, please contact the Patent Office on 0645 500505.*
- Write your answers in capital letters using black ink or you may type them.*
- If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.*
- If you have attached 'Yes' Patents Form 7/77 will need to be filed.*
- Once you have filled in the form you must remember to sign and date it.*
- For details of the fee and ways to pay please contact the Patent Office.*

ORGANIC LIGHT EMITTING MATERIAL AND DEVICE

This invention concerns new compositions and the use of such compositions in organic semi-conductor devices for example organic light emitting devices. In particular it relates to the use of luminescent boron fluoride complexes as a component of organic luminescent devices.

The use of organic electrically conducting materials as semiconductors in the fabrication of electronic devices has been explored by many investigators. Light emitting diodes have been demonstrated using a wide range of organic and molecular solids, including N,N'-diphenyl-N,N'-ditolylbenzidine, aluminium tris 8-hydroxyquinolate, 3-biphenyl-5-(4-t-butylphenyl)oxadiazole and poly(phenylene vinylene).

Conduction in organic semiconductors may be regarded as a two stage process. In the first stage, charge is injected into the organic material from the metal electrode. An example is injection of charge into evaporated thin films of 3-biphenyl-5-(4-t-butylphenyl)oxadiazole from a low work function electrode such as a magnesium metal electrode by application of a negative potential to the electrode. This injection of charge may also be regarded as an electrochemical reduction of the conducting material, or as electrochemical doping of the material. Other electronically conducting organic compounds may be subjected to charge injection by a positive potential, which is commonly applied via an electrode composed of a high work function metal such as gold. Such charge injection is commonly described as injection of positive charges denoted holes, and this description is understood to be equivalent to the extraction of electrons from the conducting material, which may also be regarded as an electrochemical doping or oxidation of the organic semiconductor.

In the second stage, charge is transported through the organic semiconductor layer. Conduction within electronically conducting organic solids involves, in practical materials, the transfer of electrical charge from one molecule to another. Such transfer of charge is described by a charge hopping or charge tunnelling mechanism which may allow an electron to overcome the energetic barrier between different molecules or molecular sub-units. In systems such as conjugated polymers including poly(acetylene) and poly(phenylenevinylene) charge can also flow along the conjugated chain by movement of

charged discontinuities in the regular bonding sequence of the polymer. In such cases, charge will normally be transferred through the bulk sample by a large number of individual molecules, and hopping or tunnelling remains an important mechanism.

In order to achieve efficient light emission from an organic electroluminescent device, it is desirable for respectively positively charged and negatively charged carriers to become simultaneously localised on a single molecule or on a small number of molecules in mutual close proximity. Such localisation of oppositely charged carriers leads to production of a molecule in its excited state, or to an excimer or exciplex comprising a plurality of molecules in an electronically excited state. Such electronically excited species, especially in the case of conjugated polymers, may also be termed exciton states.

Localisation of respectively positive and negative charge carriers may be achieved in several ways. By virtue of their opposite electronic charges, a coulombic attraction exists between them which tends to lead to localisation. Under the high electric field gradients typical of an operating device, such localisation may be ineffective in achieving high operating efficiency from devices. A known route to achieving high efficiency in devices is to use two organic semiconductor layers, one of which transports only positive carriers (holes) and the other transports only negative carriers (electrons). Under operating conditions, a high density of carriers of each sign is deposited at the interface or junction formed by the two layers, increasing the probability of their combination to form an electronically excited state. Such devices can show high efficiency but suffer the disadvantage that they require fabrication of a multilayer organic structure. Single layer structures may admit the passage of both electron and holes. Improved efficiency of operation in such single layer devices can be obtained by including in the composition a luminescent material (i.e. a dopant) at low concentration, preferably selected as having both an electron affinity higher than that of the principal negative charge carrying species in the semiconductor, and an ionisation potential lower than that of the principal positive charge carrying species in the layer. Under this condition, the luminescent material traps injected charge carriers of each polarity, increasing the probability of their combination.

The electronically excited states obtained by combination of charge carriers may be obtained in conditions of different total electronic spin, especially in so-called singlet

states and triplet states respectively. It is understood that in order to obtain high luminescence efficiency, the singlet states of excited species are most effective in many cases due to the possibility of their radiative decay to an electronic ground state with emission of a photon of light by a quantum mechanically allowed transition (fluorescence). In favourable cases, emission of light from such a singlet state occurs with high efficiency.

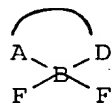
It is known that other routes exist by which the energy may be lost from an excited state, without emission of a photon of light. Non-radiative decay results in conversion of the energy to heat within the material. In most organic materials, non-radiative decay is the usual route to loss of energy from excited triplet states, as well as being a competitive process to the desirable fluorescence. A further route to loss of energy from an excited singlet state is its transition to a triplet state, followed by non-radiative decay. It is desirable in many practical devices that both direct non-radiative decay routes and the transformation of singlet states to triplet states should be minimised to achieve the highest efficiency in the device.

Some of the compounds described by the current invention are known to show high luminescence efficiency and stability, and to provide a low probability of crossing from an excited singlet state to a triplet state. In addition some of the compounds of formula I have been proposed, for example as laser dyes, and a number of compounds described by the present invention are offered commercially for this purpose.

Dodabalapur in Solid State Communications, vol 102, No2-3, pp259-267, 1997 describes the use of a pyromethene in combination with Alq3 wherein the commercially available pyromethene PM580 is present as a dopant (which may also be referred to as a light emitter) and the Alq3 is an electron transporter. In this case, no particular benefit is revealed from the use of PM580 against the known incorporation of luminescent materials such as rubrene, coumarin 6 and quinacridone derivatives into Alq3 layers in such devices. Such a route to device fabrication necessitates the use of a co-evaporation of Alq3 and dopant which requires complex facilities for its success and/or is difficult to control in practise.

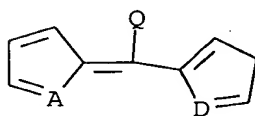
It has been unexpectedly found that compounds of the present invention show advantageous properties in respect of their ability to act as radiative combination sites for charges in organic electroluminescent structures and as electron transporting layers in OLEDs.

According to this invention an organic light emitting diode device comprises a substrate bearing an organic layer sandwiched between electrode structures wherein the organic layer comprises a hole transporter, an electron transporter and a light emitter wherein the electron transporter or the light emitter or the electron transporter and the light emitter comprise a material of general formula I:

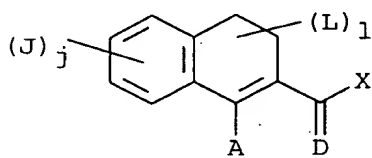


Formula I

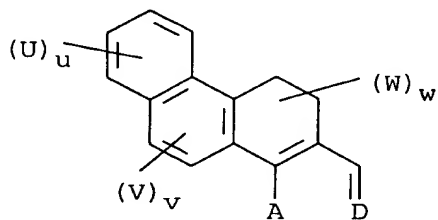
wherein $\overset{\text{A}}{\text{---}} \text{---} \overset{\text{D}}{\text{---}}$ is selected from the following:



wherein A and D are both N, and the ring systems are, independently of each other, optionally substituted with one or two or three groups independently selected from C1 - C8 straight chain or branched chain alkyl or alkoxy; Q is CN or H or C₁₋₈ straight chain or branched chain alkyl;



wherein A and D are given by O or N, X is given by C₁₋₅ straight chain or branched chain alkyl or alkoxy and the ring systems are, independently of each other, optionally substituted with one or more groups J and L independently selected from C1 - C8 straight chain or branched chain alkyl or alkoxy wherein j is selected from 0-4 and l is selected from 0-2;



wherein A and D are given by O or N and the ring systems are, independently of each other, optionally substituted with one or more groups U, V, W independently selected from C1 - C8 straight chain or branched chain alkyl or alkoxy wherein u is 0-4, v is 0-2 and w is 0-2;

provided that when the electron transport layer is Alq3 then the light emitter is not 1,3,5,7,8-pentamethyl-2,6-di-n-butylpyrromethene-difluoroborate.

According to a further aspect of this invention a composition suitable for use as an organic layer in an OLED device comprises a material of general formula I.

Compounds of Formula I show advantageous combinations of properties when used in organic light emitting devices.

The structural and other preferences are expressed below on the basis of desirable characteristics, in particular an advantageous combination of electronic orbital energy levels which is one factor determining the effectiveness of capture and recombination of electronic charge carriers in a light emitting device, luminescence quantum efficiency, solubility, processability and high chemical and photochemical stability of the material in storage and in operating devices. The present invention may give rise to devices which exhibit, inter alia, one or more of the following advantages - higher brightness, higher efficiency, purer spectral colours, longer operating life, lower cost of manufacture.

Preferably the compounds of formula I are present as:

1. light emitters
2. electron transporters and light emitters.

Preferably when the compounds of formula 1 are present only as light emitters the devices of the current invention contain a single organic layer.

Preferred materials of Formula 1 include boron fluoride adducts of pyrromethenes and of diketones, di-imines and ketoimines.

Overall preferred structures for formula I are those listed below :

- 1,2,3,5,6,7-hexamethyl-8-cyanopyrromethene-difluoroborate complex
- 1,3,5,7,8-pentamethyl-2,6-di-t-butylpyrromethene-difluoroborate
- 1,3,5,7,8-pentamethyl-2,6-di-n-butylpyrromethene-difluoroborate
- 1,3,5,7,8-pentamethyl-2,6-diethylpyrromethene-difluoroborate
- 1,3,5,7,8-pentamethylpyrromethene-difluoroborate
- 1-(1-(difluoroboryl)oxy-3,4-dihydro-naphthalen-2-yl)-ethanone
- 1-(1-(difluoroboryl)oxy-3,4-dihydro-6-methoxy-naphthalen-2-yl)-ethanone
- 1-(1-(difluoroboryl)oxy-3H-benzo(f)chromen-2-yl)-ethanone

Compounds of Formula I may be prepared by known routes, including those described in US patents 4,916,711 and 5,189,029. Typically said preparative routes comprise reaction of boron trifluoride with a diketone, diimide or ketoimide. Routes to these compounds are well known in the art.

The invention will now be described with reference to the following diagrams by way of example only:

Figure 1 illustrates an Organic Light Emitting Diode (OLED) device incorporating the materials of the present invention.

Figure 2 illustrates a matrix addressed OLED in plan view.

Figure 3 illustrates the variation of current passed as a function of applied voltage by a device given by Figure 1 - see Example 1.

Figure 4 illustrates the spectrum of emitted light corresponding to Example 1.

Figure 5 illustrates the variation of current passed as a function of applied voltage by a device given by Figure 1 - see Example 2.

Unless otherwise stated all reagents used are commercially available from the Aldrich Chemical Company.

The following compounds are illustrative examples which have been synthesised according to the present invention:

Example 1

A solution was prepared containing 0.2g N,N'-diphenyl-N,N'-di-3-tolylbenzidine purchased from Syntec GmbH of Industriepark Wolfen-Thalheim, Werkstattstrasse 188, 06766-Wolfen, Germany, 0.2g 2-(4-t-butylphenyl)-5-biphenyl-1,3,4-oxadiazole, 0.2g poly-N-vinylcarbazole and 0.00016g of 1,3,5,7,8-pentamethyl-2,6-di-n-butylpyrromethene-difluoroborate in 6.5g of 1,2-dichlorobenzene. 1,3,5,7,8-pentamethyl-2,6-di-n-butylpyrromethene-difluoroborate was purchased as laser dye PM580 from AG Electro-Optics, Farside House, Tarporley Business Centre, Tarporley, Cheshire. A 2 inch square piece of ITO coated glass (Balzers, 100 Ω /sq) was cleaned by rinsing in acetone and isopropanol, then spin coated with the above solution at 1000rpm for 30 seconds. The coated glass was immediately transferred to a heated plate at 90°C and dried for 5 minutes, then heated at 90°C in an oven for 30 minutes. The plate was transferred to a vacuum coating apparatus and aluminium was deposited onto the polymer through a perforated metal mask to form a pattern of circular aluminium contacts 3.5mm across with a thickness of 1000Å at a rate of 4 Å/second at a pressure of 10⁻⁶ torr. After coating, the device was removed from the coating equipment and electrical contact made to the ITO glass by means of a wire attached by indium solder.

A gold wire was used to make electrical contact to each of the aluminium contact pads, and a negative potential applied. Light emission could be easily observed from the ITO glass side of the device.

A Keithley 236 source measure unit was used to measure the variation of current passed by a sample device as a function of applied voltage. The results are shown in Figure 3. The spectrum of the emitted light was measured by means of a Photo Research type 714 spot photometer, and is shown in Figure 4.

Example 2

A device was prepared in the same manner and using identical conditions to those employed in Example 1, save that the metal contacts were formed from thermally evaporated samarium instead of aluminium. The variation of current passed by a sample device as a function of applied voltage is presented in Figure 5. The emission spectrum was essentially identical to that shown in Figure 4 for the device of Example 1.

Example 3

A solution was prepared containing 0.2g 2-(4-t-butylphenyl)-5-biphenyl-1,3,4-oxadiazole, 0.4g poly-N-vinylcarbazole and 0.00018g of 1,2,3,5,6,7-hexamethyl-8-cyanopyrromethene-difluoroborate complex in 6.5g of 1,2-dichlorobenzene. 1,2,3,5,6,7-hexamethyl-8-cyanopyrromethene-difluoroborate complex was purchased as laser dye PM650 from AG Electro-Optics, Farside House, Tarporley Business Centre, Tarporley, Cheshire. The solution was used in the fabrication of a light emitting device in the same manner as Example 2.

On application of a negative potential of 40V to the samarium contacts, emission of deep red light was visible.

Example 4

A substrate of ITO coated glass (Balzers, $100\Omega/\text{sq}$), was cleaned as in Example 1, and transferred to a vacuum coating apparatus. Layers of N,N'-diphenyl-N,N'-di-3-tolylbenzidine (500Å) and 1,3,5,7,8-pentamethyl-2,6-di-n-butylpyrromethene-difluoroborate (500Å) were deposited at a rate of 4 Å/second. The vacuum was broken, and a mask inserted in front of the plate. Aluminium was deposited through the mask to form an array of electrical contact pads 3.5 mm in diameter and 1000 Å thick. Contact was made to the ITO by means of a wire attached by indium solder. On application of a negative potential to the aluminium pads, emission of blue light was visible.

Example 5

A substrate of ITO coated glass (Balzers, $100\Omega/\text{sq}$), was cleaned as in Example 1, and transferred to a vacuum coating apparatus. Layers of hexamethoxy triphenylene (500Å) and 1-(1-(difluoroboryl)oxy-3,4-dihydro-naphthalen-2-yl)-ethanone (500Å) were deposited at a rate of 4 Å/second. 1-(1-(difluoroboryl)oxy-3,4-dihydro-naphthalen-2-yl)-ethanone was purchased from Synthon GmbH. The vacuum was broken, and a mask inserted in front of the plate. Aluminium was deposited through the mask to form an array of electrical contact pads 3.5 mm in diameter and 1000 Å thick. Contact was made to the ITO by means of a wire attached by indium solder. On application of a negative potential to the aluminium pads, emission of pale green light was visible.

An Organic Light Emitting Diode (OLED), which is a type of semiconductor device, suitable for incorporating materials of the present invention is illustrated in Figure 1. The device comprises two electrodes 1a, 1b, at least one of the electrodes is transparent to light of the emission wavelength of a layer of organic material 3. The other electrode may be a metal, for example Mg, Li, Ca, Al or an alloy of metals, for example MgAg, LiAl or a double metal layer, for example Li and Al or Indium Tin Oxide (ITO). One or both electrodes 1a, 1b may consist of organic conducting layers. A processing surface or substrate 2 may be made of any material which is flat enough to allow subsequent processing, for example glass, silicon, plastic. The substrate 2 may be transparent to the emitted radiation of the organic material 3. Alternatively one of the electrodes 1a, 1b may be transparent instead. Sandwiched between the electrodes 1a and 1b is a layer of organic material 3 which may itself consist of one or more layers represented here as 3a, 3b, 3c. The layer of organic material 3 possesses the following three properties: electron transporting (ET); hole transporting (HT); light emitting (LE). The materials described by the current invention are suitable for use as electron transporters and/or light emitters. If the layer of organic material 3 is a single layer then the single layer of organic material 3 must exhibit all three properties. For the case when the layer of organic material 3 is a single layer then the organic material may consist of a single material, or of two or more materials with appropriate properties mixed together. Suitable hole and/or electron transport layers include conjugated polymers such as polyphenylenevinylene (PPV) or MEHPPV, amine derivatives such as triphenylamine or TPD, heterocyclic compounds such as diphenyloxadiazole and triphenyltriazole, and heterocyclic polymers such as poly(pyridine). For the case when the layer of organic material 3 comprises more than one layer then suitable examples include:

i/ 3a = HT layer, 3b = LE layer, 3c = ET layer

ii/ 3a = HT layer, 3b = material which acts as an ET medium but also emits light, for example 1,3,5,7,8-pentamethyl-2,6-di-n-butylpyrromethene-difluoroborate

iii/ 3a = HT and LE, 3b = ET

iv/ The LE material may act as a dopant in small quantities - typically 0.5% into ET or HT or both.

Preferably in the case where the layer of organic material 3 is a multiplicity of layers, then the layer adjacent to the cathode preferentially transports electrons and/or the layer adjacent the anode preferentially transports holes. Preferably the luminescent material

has a high quantum efficiency of luminescence. The luminescent component may be combined with a charge transporting material or may be present in a separate layer.

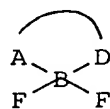
The layer of organic material 3 may be deposited on the electrode 1a by any of the following techniques: thermal evaporation under vacuum, sputtering, chemical vapour deposition, spin depositing from solution or other conventional thin film technology. The thickness of the layer of organic material 3 is typically 10-1000nm, preferably 20-200nm.

The device may contain layers 4a and 4b which are situated next to the electrodes 1a and 1b, these layers 4a and 4b may be conducting or insulating and act as a barrier to diffusion of the electrode material or as a barrier to chemical reaction at the electrode 1a, 1b and layer of organic material 3 interface. Examples of suitable materials for 4a and 4b include emeraldine which prevents indium diffusion into the layer of organic material 3 from an ITO electrode, or, for the same reason, copper phthalocyanine may be used; alternatively the addition of a thin layer (~0.5nm) of lithium or magnesium fluoride at the interface between a lithium electrode and the layer of organic material 3 may be used. The device of Figure 1 may be a single pixel device or it may be matrix addressed. An example of a matrix addressed OLED is shown in plan view in Figure 2. The display of Figure 2 has the internal structure described in Figure 1 but the substrate electrode 5 is split into strip-like rows 5l to 5m and similar column electrodes 6l to 6n, this forms an mxn matrix of addressable elements or pixels. Each pixel is formed by the intersection of a row and column electrode.

A row driver 7 supplies voltage to each row electrode 5. Similarly, a column driver 8 supplies voltage to each column electrode. Control of applied voltages is from a control logic 9 which receives power from a voltage source 10 and timing from a clock 11.

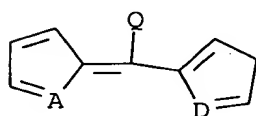
Claims

1. An organic light emitting diode device comprising a substrate bearing an organic layer sandwiched between electrode structures wherein the organic layer comprises a hole transporter, an electron transporter and a light emitter wherein the electron transporter or the light emitter or the electron transporter and the light emitter comprise a material of general formula I

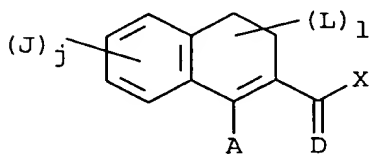


Formula I

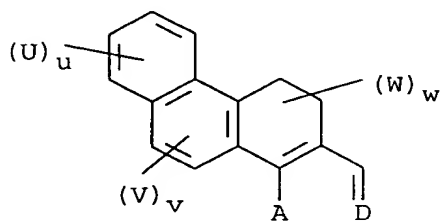
wherein $\overset{\text{A}}{\text{---}} \overset{\text{D}}{\text{---}}$ is selected from the following:



wherein A and D are both N, and the ring systems are, independently of each other, optionally substituted with one or two or three groups independently selected from C1 - C8 straight chain or branched chain alkyl or alkoxy; Q is CN or H or C₁₋₈ straight chain or branched chain alkyl;



wherein A and D are given by O or N, X is given by C₁₋₅ straight chain or branched chain alkyl or alkoxy and the ring systems are, independently of each other, optionally substituted with one or more groups J and L independently selected from C1 - C8 straight chain or branched chain alkyl or alkoxy wherein j is selected from 0-4 and l is selected from 0-2;

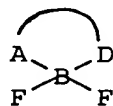


wherein A and D are given by O or N and the ring systems are, independently of each other, optionally substituted with one or more groups U, V, W independently selected from C1 - C8 straight chain or branched chain alkyl or alkoxy wherein u is 0-4, v is 0-2 and w is 0-2;

provided that when the electron transport layer is Alq3 then the light emitter is not 1,3,5,7,8-pentamethyl-2,6-di-n-butylpyromethene-difluoroborate.

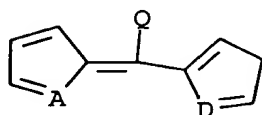
Abstract

This invention describes an organic light emitting diode device comprising a substrate bearing an organic layer sandwiched between electrode structures wherein the organic layer comprises a hole transporter, an electron transporter and a light emitter wherein the electron transporter or the light emitter or the electron transporter and the light emitter comprise a material of general formula I

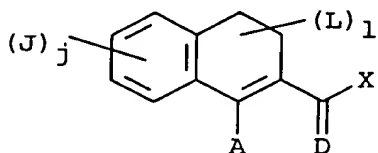


Formula I

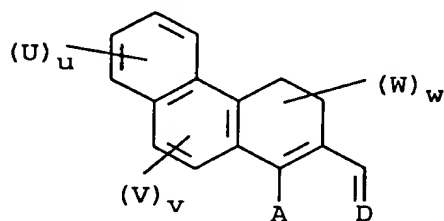
wherein A  D is selected from the following:



wherein A and D are both N, and the ring systems are, independently of each other, optionally substituted with one or two or three groups independently selected from C1 - C8 straight chain or branched chain alkyl or alkoxy; Q is CN or H or C₁₋₈ straight chain or branched chain alkyl;



wherein A and D are given by O or N, X is given by C₁₋₅ straight chain or branched chain alkyl or alkoxy and the ring systems are, independently of each other, optionally substituted with one or more groups J and L independently selected from C1 - C8 straight chain or branched chain alkyl or alkoxy wherein j is selected from 0-4 and l is selected from 0-2;



wherein A and D are given by O or N and the ring systems are, independently of each other, optionally substituted with one or more groups U, V, W independently selected from C1 - C8 straight chain or branched chain alkyl or alkoxy wherein u is 0-4, v is 0-2 and w is 0-2; provided that when the electron transport layer is Alq3 then the light emitter is not 1,3,5,7,8-pentamethyl-2,6-di-n-butylpyrromethene-difluoroborate.

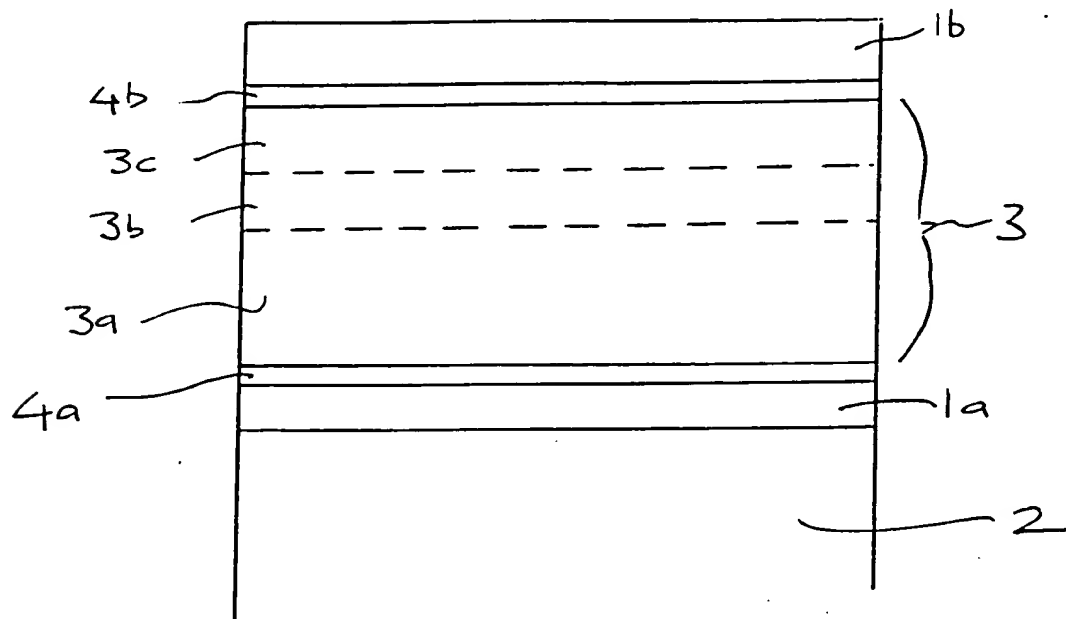


Figure 1

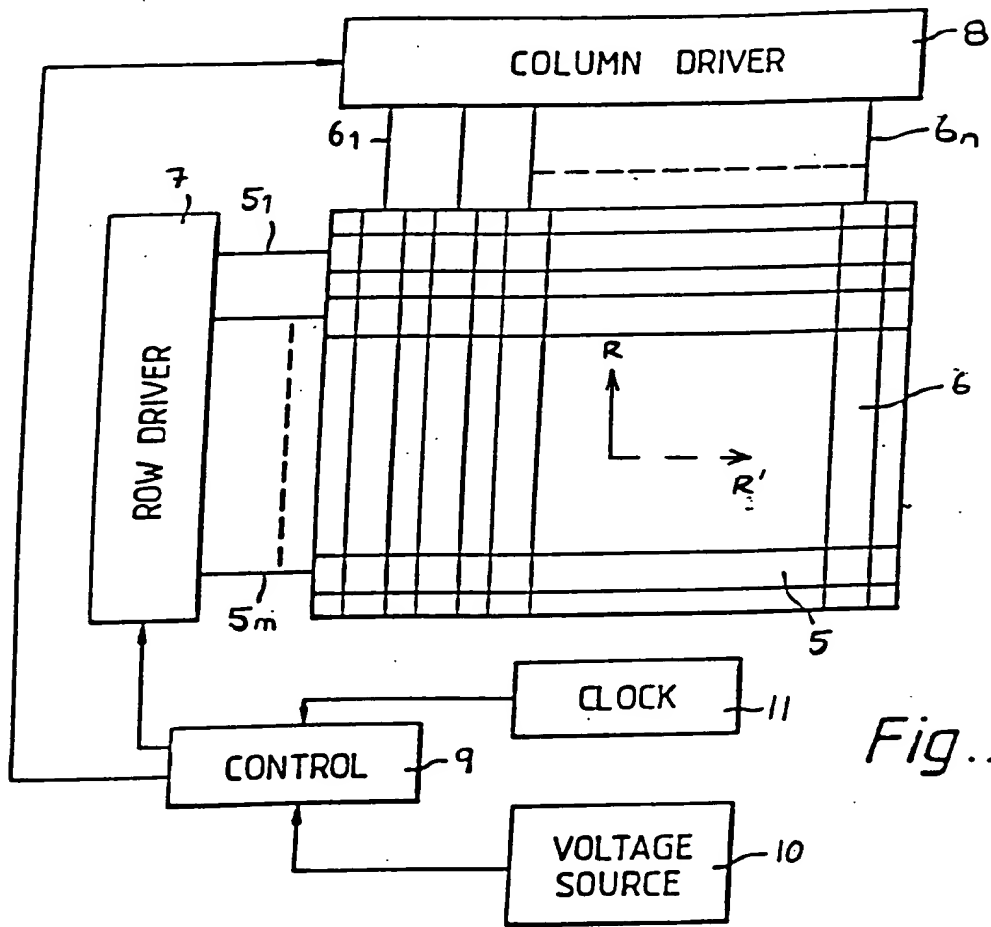


Fig. 2

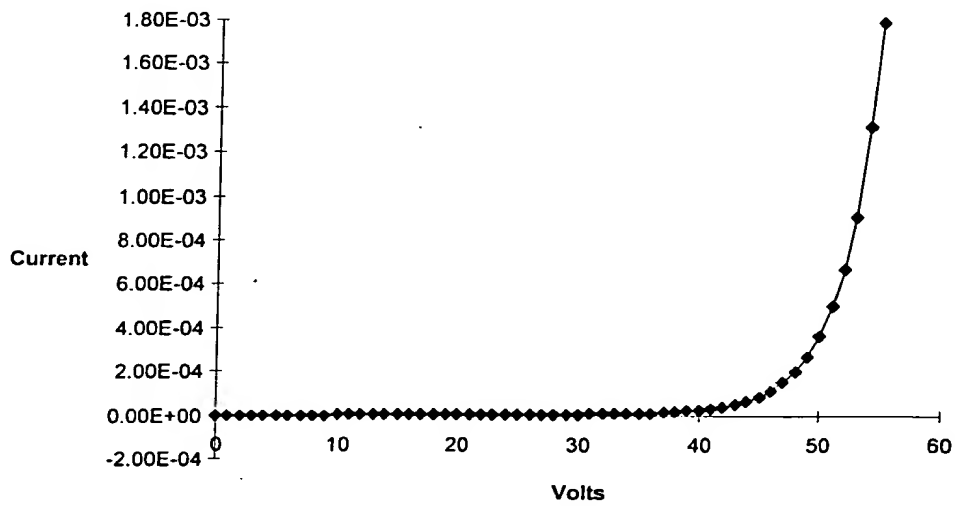


Fig 3

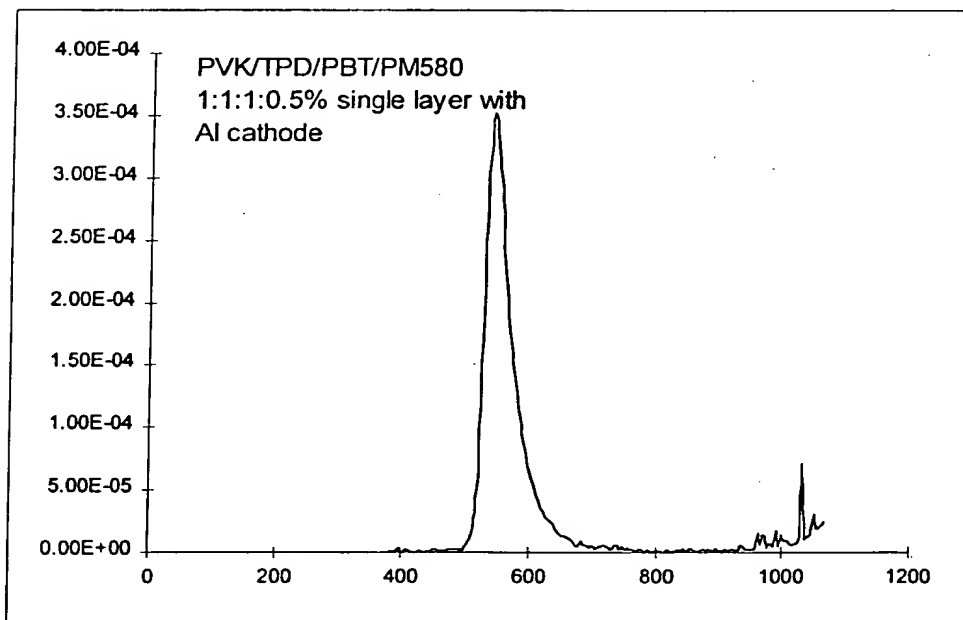


Fig 4

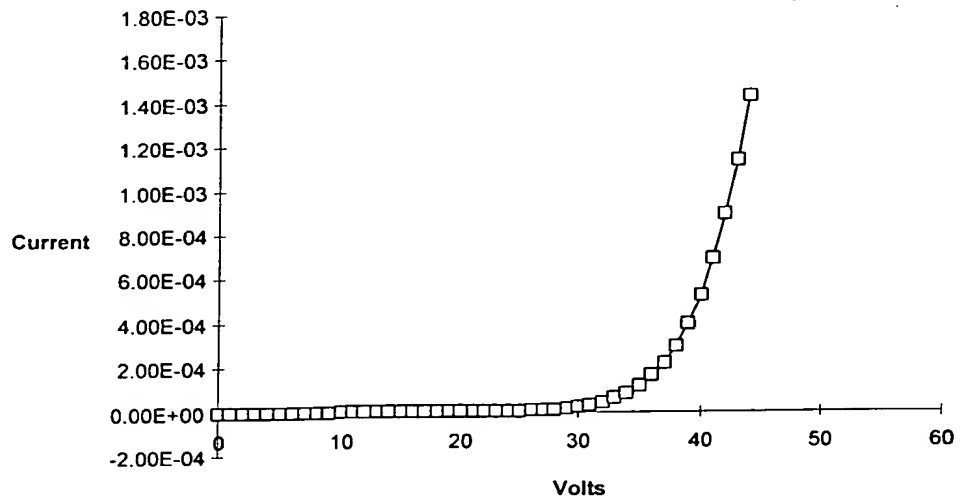


Fig 5